

APC anti-human CD194 (CCR4)

Catalog # / Size: 2397040 / 100 tests
2397035 / 25 tests

Clone: L291H4

Isotype: Mouse IgG1, κ

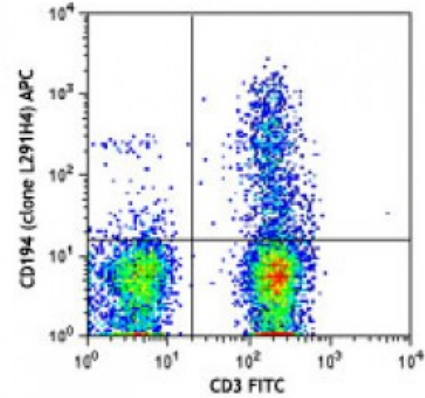
Immunogen: Human CCR4 transfected cells

Reactivity: Human

Preparation: The antibody was purified by affinity chromatography and conjugated with APC under optimal conditions. The solution is free of unconjugated APC and unconjugated antibody.

Formulation: Phosphate-buffered solution, pH 7.2, containing 0.09% sodium azide and 0.2% (w/v) BSA (origin USA).

Concentration: Lot-specific

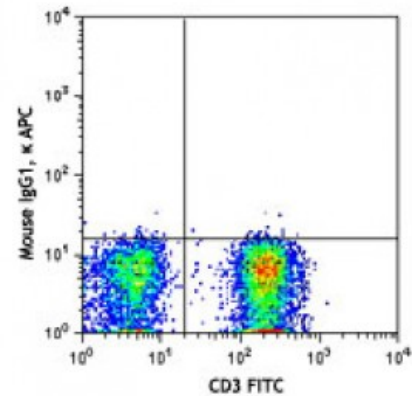


Human peripheral blood lymphocytes were stained with CD3 FITC and CD194 (clone L291H4) APC (top) or mouse IgG1, κ APC isotype control (bottom).

Applications:

Applications: Flow Cytometry

Recommended Usage: Each lot of this antibody is quality control tested by immunofluorescent staining with flow cytometric analysis. For flow cytometric staining, the suggested use of this reagent is 5 microL per million cells or 5 microL per 100 microL of whole blood. It is recommended that the reagent be titrated for optimal performance for each application.



Description: CD194, also known as CCR4, is a CC chemokine receptor. It binds CCL17 and CCL22 and is expressed on a subset of T and B cells, basophils, monocytes, and NK cells. Human Th2 cells are characterized by the expression of CCR4 and CCR8, and these receptors are regulated differently during Th2 development. Human peripheral blood Tregs can be divided into two distinct populations based on the expression of CCR4. Freshly isolated Tregs express CCR4 and presumably represent memory-type Tregs, and CCR4⁻ Tregs require CD3-mediated activation to acquire a regulatory activity. Depletion of CCR4⁺ T cells leads to Th1-type polarization of CD4⁺ T cells and augmentation of CD8⁺ T cell responses to tumor antigens. CCR4 and its ligands are important for the recruitment of memory T cells into the skin in various cutaneous immune diseases.

- Antigen References:**
1. Katschke KJ, *et al.* 2001 *Arthritis Rheum.* 44:1022.
 2. Colantonio L, *et al.* 2002 *Eur. J. Immunol.* 32:1264.
 3. Jakubzick C *et al.* 2004 *Am. J. Pathol.* 165:1211.
 4. Morimoto Y, *e*